

2018 Deafblind Census Reporting Form

Please complete and return to: Jana Villemetz - CAYSI, Arkansas Department of Education,
Special Education Unit, 1401 West Capitol, Suite 450, Little Rock, AR 72201

Complete/Update this form ONLY for individuals who have both a visual and auditory impairment. DO NOT USE for an individual with only a visual impairment.

Today's Date:

Status of this Individual's Report (please check one): ☐ DB ☐ Complex Needs ☐ Referral

Part I: Information about individual with deafblindness

Name First:

Last:

Date of Birth (MM/DD/YYYY):

/ /

Gender: ☐ Male ☐ Female

Race/Ethnicity (select the ONE that best describes the individual's race/ethnicity):

- ☐ 1 American Indian or Alaskan Native ☐ 4 Hispanic/Latino ☐ 7 Two or more races
☐ 2 Asian ☐ 5 White
☐ 3 Black /African American ☐ 6 Native Hawaiian/Pacific Islander

Living Setting (Select the ONE setting that best describes where the individual resides the majority of the year.):

- ☐ 1 Home: Birth/Adoptive Parents ☐ 5 Private Residential Facility ☐ 9 Pediatric Nursing Home
☐ 2 Home: Extended Family ☐ 6 Group Home (less than six residents) ☐ 555 Other:
☐ 3 Home: Foster Parents ☐ 7 Group Home (six or more residents)
☐ 4 State Residential Facility ☐ 8 Apartment (with non-family person(s))

Parent/Guardian Name 1 First:

Last:

Street Address:

City:

State:

ZIP Code:

Telephone (with Area Code):

County of Residence:

Parent/Guardian Name 2 First:

Last:

Street Address:

City:

State:

ZIP Code:

Telephone (with Area Code):

County of Residence:

Part II: Individual's Medical Background/Handicapping Conditions

Primary Classification of Visual Impairment (select the ONE that best describes the primary classification of the individual's visual impairment):

- ☐ 1 Low Vision ☐ 4 Totally Blind ☐ 9 Documented Functional Vision Loss
☐ 2 Legally Blind ☐ 6 Diagnosed Progressive Loss
☐ 3 Light Perception Only ☐ 7 Further Testing Needed

Cortical Vision Impairment? ☐ Yes ☐ No ☐ Unknown

Primary Classification of Hearing Impairment (select the ONE that best describes the primary classification of the individual's auditory impairment):

- ☐ 1 Mild ☐ 4 Severe ☐ 7 Further Testing Needed
☐ 2 Moderate ☐ 5 Profound ☐ 8 XXX
☐ 3 Moderately Severe ☐ 6 Diagnosed Progressive Loss ☐ 9 Documented Functional Hearing Loss

Central Auditory Processing Disorder (CAPD)?

☐ Yes ☐ No ☐ Unknown

Auditory Neuropathy?

☐ Yes ☐ No ☐ Unknown

Cochlear Implant?

☐ Yes ☐ No ☐ Unknown

Other Impairment (indicate YES or NO for each)

Physical Impairments	<input type="checkbox"/> Yes <input type="checkbox"/> No	Complex Health Care Needs	<input type="checkbox"/> Yes <input type="checkbox"/> No
Cognitive Impairments	<input type="checkbox"/> Yes <input type="checkbox"/> No	Communication Speech/Lang	<input type="checkbox"/> Yes <input type="checkbox"/> No
Behavioral Disorder	<input type="checkbox"/> Yes <input type="checkbox"/> No	Other: _____	<input type="checkbox"/> Yes <input type="checkbox"/> No

Etiology (please indicate the ONE etiology from the lists below that best describes the primary etiology of the individual's primary disability. Please indicate "Other" if none of the listed etiologies are the primary disability):

Hereditary/Chromosomal Syndromes and Disorders

- | | |
|---|---|
| <input type="radio"/> 101 Aicardi syndrome | <input type="radio"/> 130 Marshall syndrome |
| <input type="radio"/> 102 Alport syndrome | <input type="radio"/> 131 Maroteaux-Lamy syndrome (MPS VI) |
| <input type="radio"/> 103 Alstrom syndrome | <input type="radio"/> 132 Moebius syndrome |
| <input type="radio"/> 104 Apert syndrome (Acrocephalosyndactyly, Type!) | <input type="radio"/> 133 Monosomy Tenp |
| <input type="radio"/> 105 Bardet-Biedl syndrome (Laurence Moon-Biedl) | <input type="radio"/> 134 Morquio syndrome (MPS IV-B) |
| <input type="radio"/> 106 Batten disease | <input type="radio"/> 135 NF One - Neurofibromatosis (Von Recklinghausen) |
| <input type="radio"/> 107 CHARGE association | <input type="radio"/> 136 NF Two- Bilateral Acoustic Neurofibromatosis |
| <input type="radio"/> 108 Chromosome eighteen, Ring eighteen | <input type="radio"/> 137 Norrie disease |
| <input type="radio"/> 109 Cockayne syndrome | <input type="radio"/> 138 Optico-Cochleo-Dentate Degeneration |
| <input type="radio"/> 110 Cogan syndrome | <input type="radio"/> 139 Pfeiffer syndrome |
| <input type="radio"/> 111 Cornelia de Lange | <input type="radio"/> 140 Prader-Willi |
| <input type="radio"/> 112 Cri du chat syndrome (Chromosome 5p-Syndrome) | <input type="radio"/> 141 Pierre-Robin syndrome |
| <input type="radio"/> 113 Crigler-Najjar syndrome | <input type="radio"/> 142 Refsum syndrome |
| <input type="radio"/> 114 Crouzon syndrome (Craniofacial Dysostosis) | <input type="radio"/> 143 Scheie syndrome (MPS I-S) |
| <input type="radio"/> 115 Dandy Walker syndrome | <input type="radio"/> 144 Smith-Lemli-Optiz (SLO) syndrome |
| <input type="radio"/> 116 Down syndrome (Trisomy Twenty-one) | <input type="radio"/> 145 Stickler syndrome |
| <input type="radio"/> 117 Goldenhar syndrome | <input type="radio"/> 146 Sturge-Weber syndrome |
| <input type="radio"/> 118 Hand-Schuller-Christian (Histiocytosis X) | <input type="radio"/> 147 Treacher Collins syndrome |
| <input type="radio"/> 119 Hallgren syndrome | <input type="radio"/> 148 Trisomy thirteen (Patau syndrome) |
| <input type="radio"/> 120 Herpes-Zoster (or Hunt) | <input type="radio"/> 149 Trisomy eighteen (Edwards syndrome) |
| <input type="radio"/> 121 Hunter syndrome (MPSII) | <input type="radio"/> 150 Turner syndrome |
| <input type="radio"/> 122 Hurler syndrome (MPS I-H) | <input type="radio"/> 151 Usher I syndrome |
| <input type="radio"/> 123 Kearns-Sayre syndrome | <input type="radio"/> 152 Usher II syndrome |
| <input type="radio"/> 124 Klippel-Feil sequence | <input type="radio"/> 153 Usher III syndrome |
| <input type="radio"/> 125 Klippel-Trenaunay-Weber syndrome | <input type="radio"/> 154 Vogt-Koyanagi-Harada syndrome |
| <input type="radio"/> 126 Kniest Dysplasia | <input type="radio"/> 155 Waardenburg syndrome |
| <input type="radio"/> 127 Leber congenital amaurosis | <input type="radio"/> 156 Wildervanck syndrome |
| <input type="radio"/> 128 Leigh disease | <input type="radio"/> 157 Wolf-Hirschhorn syndrome (Trisomy 4p) |
| <input type="radio"/> 129 Marfan syndrome | <input type="radio"/> 199 Other |

OR Pre-Natal/Congenital Complications

- | | | |
|---|--|---|
| <input type="radio"/> 201 Congenital Rubella Syndrome | <input type="radio"/> 205 Fetal Alcohol Syndrome | <input type="radio"/> 209 Neonatal Herpes Simples (HSV) |
| <input type="radio"/> 202 Congenital Syphilis | <input type="radio"/> 206 Hydrocephaly | <input type="radio"/> 299 Other |
| <input type="radio"/> 203 Congenital Toxoplasmosis | <input type="radio"/> 207 Maternal Drug Use | |
| <input type="radio"/> 204 Cytomegalovirus (CMV) | <input type="radio"/> 208 Microcephaly | |

OR Post-Natal/Non-Congenital Complications

- | | | | |
|---|--|--|---------------------------------|
| <input type="radio"/> 301 Asphyxia | <input type="radio"/> 304 Infections | <input type="radio"/> 307 Stroke | <input type="radio"/> 399 Other |
| <input type="radio"/> 302 Direct Trauma to the eye and/or ear | <input type="radio"/> 305 Meningitis | <input type="radio"/> 308 Tumors | |
| <input type="radio"/> 303 Encephalitis | <input type="radio"/> 306 Severe Head Injury | <input type="radio"/> 309 Chemically Induced | |

OR Related to Prematurity☐ 401 Complications of Prematurity**Undiagnosed**☐ 501 No Determination of Etiology

Part III: IDEA

Funding Category (please indicate the funding category under which the individual was receiving services on **December 1, 2018**):

- ☐ 1 IDEA Part B (three through twenty-one years) ☐ 3 Not reported under Part B or Part C
☐ 2 IDEA Part C (birth through two years)

-----Part C-----

Special Education Status/Part C Exiting (please indicate the **ONE** code that best describes the individual's special education program status):

- ☐ 0 In a Part C early intervention program ☐ 6 Deceased
☐ 1 Completion of IFSP prior to reaching max age for Pt C ☐ 7 Moved out of state
☐ 2 Eligible for IDEA, Part B ☐ 8 Withdrawal by parent/guardian
☐ 3 Not eligible for Pt B, referral to other program ☐ 9 Attempts to reach parent and/or child unsuccessful
☐ 4 Not eligible for Pt B, exit w/no referrals
☐ 5 Part B eligibility not determined

Part C Category Code (please indicate the primary category code under which the individual was reported on the Part C, IDEA Child Count - select only ONE. Note: Preschoolers who turned 3 years old during the reporting period (Dec 1, 2017 -Dec 1, 2018) and who have transitioned to Part B services may also be reported under Part B-Exiting status:2018

- ☐ 1 At-risk ☐ 9 Other Health Impairment
☐ 2 Developmentally Delayed ☐ 10 Specific Learning Disability
☐ 3 Mental Retardation ☐ 11 Deaf-blindness
☐ 4 Hearing Impairment (includes deafness) ☐ 12 Multiple Disabilities
☐ 5 Speech or Language Impairment ☐ 13 Autism
☐ 6 Visual Impairment (includes blindness) ☐ 14 Traumatic Brain Injury
☐ 7 Emotional Disturbance ☐ 888 Not Reported under Part C of IDEA
☐ 8 Orthopedic Impairment

-----Part B-----

Special Education Status/Part B Exiting (please indicate the **ONE** code that best describes the individual's special education program status):

- ☐ 0 In ECSE or school-aged Special Education Program ☐ 5 Died
☐ 1 Transferred to regular education ☐ 6 Moved, Known to be Continuing
☐ 2 Graduated with regular diploma ☐ 7 (intentionally not used)
☐ 3 Received a certificate ☐ 8 Dropped out
☐ 4 Reached Maximum Age

Part B Category Code (please indicate the primary category code under which the individual was reported on the Part B, IDEA Child Count Note: Preschoolers who turned 3 years old during the reporting period (December 1, 2017-Dec 1, 2018) and who have transitioned to Part C early intervention services may also be reported under Part C-Exiting status: 2018

- ☐ 1 Intellectual Disability ☐ 9 Deaf-blindness
☐ 2 Hearing Impairment (includes deafness) ☐ 10 Multiple Disabilities
☐ 3 Speech or Language Impairment ☐ 11 Autism
☐ 4 Visual Impairment (includes blindness) ☐ 12 Traumatic Brain Injury
☐ 5 Emotional Disturbance ☐ 13 Developmentally Delayed - age 3 through 9
☐ 6 Orthopedic Impairment ☐ 14 Non-Categorical
☐ 7 Other Health Impairment ☐ 15 Not Reported under Part B of IDEA
☐ 8 Specific Learning Disability

Deaf-Blind Project Exiting Status:

- ☐ 0 Eligible to receive services from DB Project
☐ 1 No longer eligible to receive services from DB Project

Participation in Statewide Assessments

- ☐ 1 Regular grade-level state assessment
☐ 2 Regular grade-level state assessment w/accommodations
☐ 3 Alternate assessment
☐ 4 XXXX
☐ 5 XXXX
☐ 6 Not yet required at age or grade level

Educational Setting (indicate the ONE educational setting code from the appropriate age subcategory that best describes the individual's educational setting. Please specify "Other" if none of the provided codes apply):

Early Intervention Setting

Birth through 2 years of age (if the individual is in this category, please check the ONE box indicating the service(s) setting).

- ☐ 1 Home
☐ 2 Community-based settings
☐ 3 Other Settings

ECSE (3-5) Settings

- ☐ 1 In a regular EC program 10+ hrs/wk with services
☐ 2 In a regular EC program 10+ hrs/wk services elsewhere
☐ 3 In a regular EC program less than 10 hrs/wk with services
☐ 4 In a regular EC program less than 10 hrs/week-services elsewhere
☐ 5 Attending a separate class
☐ 6 Attending a separate school
☐ 7 Attending a residential facility
☐ 8 Service provider location
☐ 9 Home

School aged (6-21) Settings

- ☐ 10 Inside the regular class 80% or more of the day
☐ 11 Inside the regular class 40% to 79% of day
☐ 12 Inside the regular class less than 40% of the day
☐ 13 Separate School
☐ 14 Residential facility
☐ 15 Homebound/Hospital
☐ 16 Correctional facilities
☐ 17 Parentally placed in private schools

Assistive Technology

Corrective Lenses ☐ Yes ☐ No ☐ Unknown

Assistive Listening Devices ☐ Yes ☐ No ☐ Unknown

Additional Assistive Technology ☐ Yes ☐ No ☐ Unknown

Intervener Services OR 1:1 Paraprofessional

Intervener ☐ Yes ☐ No ☐ Unknown Has a 1:1 ☐ Yes ☐ No ☐ Unknown

School Information

Agency OR School:

Street Address:

City: State: ZIP Code:

Telephone Number: Fax Number:

Teacher's Name:

Teacher's Email:

School District:

LEA Supervisor: Date:

Please return this form by February 11, 2019 to:

Jana Villemetz, Arkansas CAYSI Project-Children And Youth with Sensory Impairment and other disabilities

ADE, Special Education Unit
 1401 West Capitol, Suite 450
 Little Rock, AR 72201

Telephone: 501-682-4222; Fax: 501-682-4248

Thank you for completing this. It will assist in program development and funding.